

Biomechanical simulations and 3D printing for endovascular device testing

Michele Conti, Stefania Marconi, Ferdinando Auricchio

DICAr Department, University of Pavia, Pavia, Italy

Abstract

Endovascular aortic repair is a minimally invasive procedure to treat aortic diseases such as aneurysms and dissections. Thanks to technological advancements, such procedure has steadily shifted from the abdominal aorta towards the ascending part, *i.e.*, near the heart, calling for an extensive and comprehensive benchmarking of (novel) endografts. Given such considerations, we have exploited porcine aorta with a pulse duplicator to analyse the mechanical interaction between the endovascular device and the native tissue.

Our results have implications for using the porcine aorta as a model for human aorta in research. Particularly, the combination of *in vitro* tests performed using *ex-vivo* tissue, integrated validated patient-specific numerical simulations, mock arteries manufactured by 3D printing, can offer important insight on biomechanical impact of endograft design on post-operative aortic mechanical response.

Introduction

Endovascular aortic repair (EVAR) is a minimally invasive procedure to treat aortic diseases such as aneurysms and dissections. Thanks to technological advancements, such a procedure has steadily shifted from the abdominal aorta towards the ascending part, *i.e.*, near the heart, calling for an extensive and comprehensive benchmarking of (novel) endograft. Given such considerations, we have exploited porcine aorta with a pulse duplicator to analyse the mechanical interaction between the endovascular device and the native tissue.

Materials and Methods

We have investigated the effect of thoracic endovascular aortic repair (TEVAR) on aortic stiffness by measuring aortic pulse wave velocity (PWV) in an *ex vivo* porcine model.¹ In particular, fifteen fresh porcine thoracic aortas were connected to a benchtop pulsatile. Intraluminal pressures were recorded in the ascending aorta and at the celiac trunk using a needle connected to a pressure sensor. The distance between the needles was divided by the time difference between the base of the pressure peaks to calculate aortic PWV at baseline and after stent-graft deployment and distal stent-graft extension (Figure 1).

Similarly, twenty fresh thoracic porcine aortas were connected to a mock circulatory loop driven by a centrifugal flow pump. A high definition camera captured diameters at five different pressure levels (100, 120, 140, 160, and 180 mmHg), before and after TEVAR,² as function of different degree of oversizing. Moreover, we have consistently and systematically compared published data on porcine and human thoracic aortic stiffness from different studies.³

Results

In our *ex-vivo* experimental setup, aortic stiffness increased after stent-graft deployment, dependent on the percentage of the aorta that was covered by stent graft. TEVAR stiffened the thoracic aorta by 2-

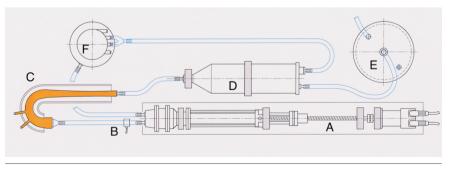


Figure 1. Schematic representation of the pulsatile system. A, pump; B, flow meter; C, exvivo porcine aortic specimen; D, resistance chamber; E, windkessel compartment; F, water reservoir.

Correspondence: Michele Conti, DICAr Department, University of Pavia, Pavia, Italy. E-mail: michele.conti@unipv.it

Key words: Aortic model; *ex vivo* experiments; pulse-duplicator; 3d-printing.

Acknowledgements: the authors acknowledge the contribution of the research group lead by Prof. S. Trimarchi (University of Milan) and Prof. F. Moll (UMCU-Utrecht-NL).

Conference presentation: this paper was presented at the Second Centro 3R Annual Meeting - 3Rs in Italian Universities, 2019, June 20-21, University of Genoa, Italy.

Received for publication: 28 October 2019. Accepted for publication: 6 November 2019.

This work is licensed under a Creative Commons Attribution NonCommercial 4.0 License (CC BY-NC 4.0).

©Copyright: the Author(s), 2019 Licensee PAGEPress, Italy Biomedical Science and Engineering 2019; 3(s2):86 doi:10.4081/bse.2019.86

fold. Such segmental stiffening may diminish the Windkessel function considerably and might be associated with TEVAR-related complications, including stent-graftinduced dissection and aneurysmal dilatation. Furthermore, our results show that the stiffness of young porcine aortas is similar to that of human tissue aged under 60 years and less stiff than human tissue aged 60 years or more.

Conclusions

Our results have implications for using the porcine aorta as a model for human aorta in research. In particular, the combination of *in vitro* tests performed using *ex-vivo* tissue, integrated validated patient-specific numerical simulations,⁴ mock arteries manufactured by 3D printing,⁵ can offer important insight on biomechanical impact of endograft design on post-operative aortic mechanical response.

References

- de Beaufort HW, Conti M, Kamman AV, et al. Stent-graft deployment increases aortic stiffness in an ex vivo porcine model. Ann Vasc Surg 2017;43:302-8.
- 2. Nauta FJ, de Beaufort HW, Conti M, et al. Impact of thoracic endovascular aor-





tic repair on radial strain in an ex vivo porcine model. Eur J Cardiothorac Surg 2017;51:783-9.

3. de Beaufort HW, Ferrara A, Conti M, et al. Comparative analysis of porcine and human thoracic aortic stiffness. Eur J

Vasc Endovasc Surg 2018;55:560-6.

4. Romarowski RM, Conti M, Morganti S, et al. Computational simulation of TEVAR in the ascending aorta for optimal endograft selection: A patient-specific case study. Comput Biol Med 2018;103:140-7.

 Marconi S, Lanzarone E, van Bogerijen GH, et al. A compliant aortic model for in vitro simulations: Design and manufacturing process. Med Eng Phys 2018;59:21-9.